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Key tumor suppressor genes inactivated by “greater promoter” methylation and somatic mutations in head and neck cancer

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Keywords: Head and Neck Squamous Cell Carcinoma, Tumor Suppressor Genes, DNA methylation, somatic mutations, integration analysis

Tumor suppressor genes (TSGs) are commonly inactivated by somatic mutation and/or promoter methylation; yet, recent high-throughput genomic studies have not identified key TSGs inactivated by both mechanisms. We pursued an integrated molecular analysis based on methylation binding domain sequencing (MBD-seq), 450K Methylation arrays, whole exome sequencing, and whole genome gene expression arrays in primary head and neck squamous cell carcinoma (HNSCC) tumors and matched uvulopalatopharyngoplasty tissue samples (UPPPs). We uncovered 186 down-regulated genes harboring cancer specific promoter methylation including *PAX1* and *PAX5* and we identified 10 key tumor suppressor genes (*GABRB3*, *HOXC12*, *PARP15*, *SLCO4C1*, *CDKN2A*, *PAX1*, *PIK3AP1*, *HOXC6*, *PLCB1*, and *ZIC4*) inactivated by both promoter methylation and/or somatic mutation. Among the novel tumor suppressor genes discovered with dual mechanisms of inactivation, we found a high frequency of genomic and epigenomic alterations in the *PAX* gene family of transcription factors, which selectively impact canonical *NOTCH* and *TP53* pathways to determine cell fate, cell survival, and genome maintenance. Our results highlight the importance of assessing TSGs at the genomic and epigenomic level to identify key pathways in HNSCC, deregulated by simultaneous promoter methylation and somatic mutations.

Introduction

Head and neck squamous cell carcinoma (HNSCC) is the sixth most common cancer worldwide with an approximate 50% five-year survival rate.¹ Two groups that independently studied the genetic origins of HNSCC reported inactivating mutations in *NOTCH1*.^{1,2} This was the first strong evidence of *NOTCH1* mutations in solid tumors; analysis of the mutations suggested that *NOTCH1* might act as a tumor suppressor gene (TSG) in HNSCC.^{1,2} Notwithstanding this important finding, and contrary to original expectations, these and other detailed analyses of HNSCC did not uncover a great number of recurrent somatic mutations in novel genes.^{3,4}

The number of known mutations and specific mutational hotspots in HNSCC tumors only partially explains their biological complexity and limits the development of novel diagnostic markers and therapeutic agents. *TP53* was again identified as the most commonly mutated gene in HNSCC and, while mutant *TP53* has been associated with poor survival,⁵ the most important biologic consequences of this alteration have been elusive. Moreover, it was also known that overall and disease-specific survival is higher in patients with HPV-associated HNSCC tumors,⁶ and that this distinct molecular and pathologic subtype displays an average of 4 somatic mutations per tumor, while HPV-negative HNSCC tumors harbor 20.¹ HPV-associated HNSCCs have a distinctly different molecular landscape from HPV-negative HNSCCs. *TP53* is rarely identified as mutated in HPV-positive HNSCC patients because the HPV E6 oncoprotein silences

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TP53, protecting the cells from apoptosis and senescence, while the HPV E7 oncoprotein deregulates the cell cycle.^{7,8} *CDKN2A*, a principal cyclin-dependent kinase inhibitor that decelerates the cell cycle, is lost in HPV-negative HNSCC⁹ and amplified in HPV-positive HNSCC.¹⁰ HNSCCs also exhibit many chromosomal abnormalities, including amplifications of the 11q13 region containing the *cyclin D1* gene and the 7p11 region encoding *EGFR*, which lead to proto-oncogene activation.¹

Aberrant DNA methylation of CpGs in the proximity of predicted transcription start sites (TSS) often leads to alterations in gene function and pathway deregulation in human cancer. Epigenetic events linked to TSG inactivation through promoter methylation, are more frequent events than somatic mutations in cancer, and may be driving tumorigenic initiation and progression. Promoter methylation of *CDK2NA*, *HOXA9*, *NID2*, *EDNRB*, *KIF1A*, and *DCC* have previously been identified and characterized in HNSCC.¹¹⁻¹³

To date, recent high-throughput methylation studies^{14,15} have not focused on the relative contribution of TSGs inactivation by DNA methylation and somatic mutations in oncogenesis and may have severely underestimated the true frequency of inactivation of key TSGs and signaling pathways. We tested the hypothesis that TSG promoter methylation predominantly occurs in genes or pathways with well-known somatic mutations and/or deletions in most HNSCC tumors, including *TP53*, *CDK2NA*,¹⁶ and, more recently, *NOTCH1*^{1,2} and *FAT1*,¹⁷ as well as in recently described genes with low frequency mutations.¹⁸

Results

Patient selection

One hundred and eight (108) head and neck squamous cell carcinoma (HNSCC) and 35 uvulopalatopharyngealplasty (UPPP) patients were consented for this study at Johns Hopkins Medical Institutions hospitals and MD Anderson Cancer Center. The study was approved by the Ethics Committee of each participating hospital, as well as by the Johns Hopkins Institutional Review Board. The 143 samples were divided into Discovery and Prevalence cohorts. The Discovery cohort consisted of 32 HNSCC and 16 UPPP samples. The Prevalence cohort consisted of 76 HNSCC and 19 UPPP samples. The 32 tumor samples selected for the Discovery cohort were accrued in Hopkins (n=17) and in MD Anderson (n=15). The 35 UPPP samples were accrued at Hopkins.

Johns Hopkins component

Fresh-frozen surgically resected HNSCC (*n* = 93) and UPPP (*n* = 35) tissue samples were obtained from patients at Johns Hopkins Medical Institutions in Baltimore. HNSCC tissue was analyzed by frozen section histology to estimate neoplastic cellularity. In order to enrich the samples for neoplastic cells, normal tissue was removed from the samples using macro-dissection based on the frozen section histology. HPV tumor status was determined for oropharyngeal tumors per standard clinical care using in situ hybridization. Hybridization was performed using the HPV III Family16 probe set that captures HPV genotypes

16, 18, 33, 35, 45, 51, 52, 56, and 66. HPV16-positive controls included an HPV16-positive oropharyngeal cancer and the SiHa and CaSki cell lines. HPV tumor status was also determined by E6/E7 PCR primer amplification.

MD Anderson component

Fresh-frozen surgically resected tumor samples (*n* = 15) were obtained from consented patients treated for HNSCC at the University of Texas MD Anderson Cancer Center, under an Institutional Review Board approved protocol. Frozen tissue was embedded in optimal cutting temperature compound and cryosections from the top and middle of specimens were stained with hematoxylin and eosin prior to being evaluated by a pathologist for the presence of >60% tumor nuclei content or absence of tumor (i.e., normal). Samples that passed this criterion were sectioned all the way through and washed once in PBS prior to isolating genomic DNA using an ArchivePure DNA purification kit.

Table 1 describes the clinical attributes of the combined 143 patient samples from the Discovery and Validation cohorts, as well as of the 289 samples from The Cancer Genome Atlas that were used to confirm the *TP53*mut/*PAX5* met and the *NOTCH1*mut/*PAX1*met associations. The 32 HNSCC patient samples selected for Discovery in this epigenetic study were the same used by Agrawal et al. for the genetic study of HNSCC¹. This study design allowed us to analyze mutation, methylation, and expression data from the same HNSCC patients. We queried the methylome of these 32 HNSCC patients and frequency matched 16 UPPP normal controls using the Human Methylation 450K Beadchip (Illumina). However, the HNSCC transcriptome was queried with different platforms. In Hopkins the GeneST1.0 arrays (Affymetrix) were used to query the HNSCC and UPPP transcriptome and in MD Anderson they queried the HNSCC transcriptome with the Human Exon 1.0ST arrays (Affymetrix). The integrated methylation/mRNA expression analysis we are reporting was performed using transcriptome data from the 32 samples (16 HNSCC and 16 UPPP) accrued at Hopkins because we did not find a satisfactory method of combining GeneST1.0 and Human Exon 1.0ST array data. Nor did we have transcriptome and methylome data from normal patient samples collected at MD Anderson to compare with the results obtained with patients accrued at Hopkins.

Characterization of the HNSCC methylome using MBD-seq

In the first part of our study, we used a methylated DNA binding domain based sequencing (MBD-seq) approach similar, in principle, to what has been previously described,¹⁹ but with significant modifications (Fig. 1). This analysis was performed on a subset of tumors from the same Discovery Cohort in which Agrawal et al. discovered and mapped mutations in HNSCC,¹ comprising ten patients and ten frequency matched normal controls (uvulopalatopharyngoplasty tissue samples [UPPP]). The ten tumor samples were obtained before chemotherapy or radiation treatment, ensuring that the changes we identified are truly reflective of tumor biology, and were micro-dissected to achieve a neoplastic cellularity of greater than 60%. Following MBD-seq, an average of 48.9 million 50 bp reads was obtained for each

Table 1. Summary of clinical attributes of samples in Discovery, Prevalence, and TCGA cohorts

	Discovery			Prevalence			TCGA	
	Normal	HPV+	HPV-	Normal	HPV+	HPV-	HPV+	HPV-
	(n = 16)	(n = 3)	(n = 29)	(n = 19)	(n = 4)	(n = 72)	(n = 35)	(n = 244)
Sex								
Male	9(56%)	3(100%)	18(62%)	14(74%)	3(75%)	52(72%)	31(89%)	172(70%)
Race								
Non-Latino White	9(56%)	3(100%)	22(76%)	11(58%)	3(75%)	62(86%)	33(94%)	209(86%)
African American/ Black	7(44%)	0	4(14%)	7(37%)	1(25%)	7(10%)	2(6%)	24(10%)
Other	0	0	3(10%)	1(5%)	0	3(4%)	0	11(5%)
Smoking								
Yes	0	1(33%)	23(79%)	0	2(50%)	26(38%)	25(75%)	195(80%)
No	16(100%)	2(67%)	6(21%)	19(100%)	2(50%)	12(18%)	10(29%)	41(17%)
Unknown	0	0	0		0	34(47%)	0	8(3%)
Tumor Site								
Oral Cavity		0	25(86%)		0	35(49%)	12(34%)	160(66%)
Oropharynx		3(100%)	2(7%)		3(75%)	21(29%)	21(60%)	12(5%)
Larynx		0	2(7%)		0	10(14%)	1(3%)	71(29%)
Hypopharynx		0	0		0	4(5%)	1(3%)	1(0.4%)
Unknown		0	0		1(25%)	2(3%)	0	0
T stage								
I		0	1(4%)		0	0	3(9%)	17(7%)
II		3(100%)	6(21%)		0	1(1%)	10(29%)	63(26%)
III		0	12(41%)		0	2(3%)	2(6%)	54(22%)
IV		0	9(31%)		4(100%)	63(88%)	9(26%)	87(36%)
Unknown		0	1(4%)		0	6(8%)	11(31%)	23(9%)

sample, with an average of 68% of these reads aligning to the hg19 genome, 67% of which were aligning uniquely (Data set 1).

Our purpose was to study alterations in the promoter region of TSGs and for this study we focused on the “greater promoter,” a region that encompasses the well-studied proximal promoter that harbors CpG Islands (1500 bases up and downstream from the TSS) and the distal promoter (6000 bp upstream of the TSS), which includes recently identified CpG Island Shores²⁰ and Shelves (Fig. 2A).²¹ (From this point on, wherever the term promoter is used it refers to the greater promoter region as defined here).

We used two independent and highly validated analytical approaches, model-based analysis of ChIP-seq (MACS)²² and Bump hunting,²³ to identify methylation changes across the HNSCC genome. MACS identified 9648 alterations, 60% of which were gains in methylation events (Fig. 2B). The vast majority of the methylome changes identified by MACS (73%) were observed outside of the promoter region. Within the promoter region, we observed mostly (89%) gain of methylation events. The majority of the methylation loss identified by MACS (93%) occurred outside the promoter region. These genome-wide methylation motifs were integrated with the differentially methylated

regions (DMRs) identified by Bump hunting to obtain the first detailed next-gen analysis of the HNSCC promoter methylome (Data set 2; Tables S1 and S2).

Confirmation of methylated sites using 450K arrays and integration with expression arrays

To validate the MBD-seq results, we evaluated genome-wide differential methylation with the Infinium HumanMethylation450K Beadchip (450K array) in 41 cancer (including the 10 samples sequenced with MBD-seq) and 16 UPPP samples. The intersection of genome-wide methylation sequencing and methylation array screens uncovered 316 genes, which harbor promoter methylation in HNSCC (Table S3). To determine the extent of correlation between differential methylation and mRNA expression patterns, we performed mRNA expression microarray analysis (Affymetrix Gene ST 2.0) using 16 tumor and 16 UPPP samples from the samples included in the cohort analyzed with the 450K array platform. We found close to 60% concordance between concurrent greater promoter methylation and gene downregulation. 186 methylated genes were found to harbor methylation and downregulation; *PAX1* and *PAX5* exhibited the greatest expression loss (Fig. 3A; Table S4A). Table S4B shows the relationship between methylation frequency

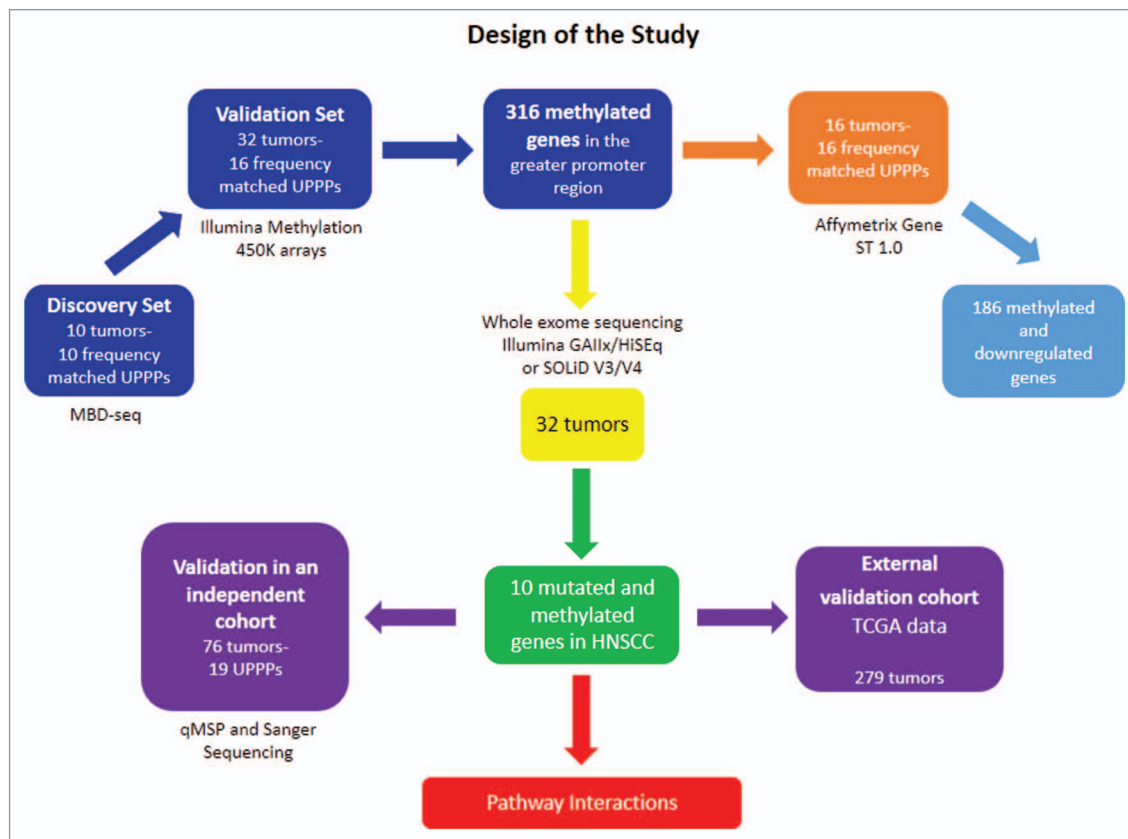


Figure 1. Representation of the workflow of the study. The figure ascribes all the platforms and techniques used in the discovery and the two independent validation sets. The number of samples recruited its time are also depicted in brackets.

of the significantly downregulated candidate genes with $FC < -2$ and the expression levels of these candidate genes for the 16 tumor samples queried with mRNA arrays in the Discovery cohort. We have also included **Figure S1** to show the expression level box plots for these candidate genes in the 16 HNSCC and 16 UPPP samples queried in the Discovery cohort.

Integration of promoter methylation with somatic mutation profiles

We subsequently intersected the promoter methylome with the mutational landscape of HNSCC¹ and identified concurrent promoter methylation and somatic mutations in ten tumor suppressor genes: *GABRB3*, *HOXC12*, *PARP15*, *SLCO4C1*, *CDKN2A*, *PAX1*, *PIK3API*, *HOXC6*, *PLCB1*, and *ZIC4* (**Fig. 3B**). *CDKN2A* (p16), one of the most frequently altered tumor suppressor genes in human cancer by mutation, methylation, and/or deletion was confirmed on this list. The rest of the genes displayed a very low mutation frequency in HNSCC, together with frequent inactivation by promoter methylation.

Unbiased genome-wide and gene set enrichment analyses

Unbiased genome-wide analyses were performed to visualize and interpret the large amount of data produced by the sequencing and microarray experiments. Unsupervised hierarchical clustering of the differential methylation events in HNSCC revealed a genome-wide loss of methylation. More than half of the genes in cancer have lost methylation when compared with normal samples at the genome-wide level. This massive loss of

methylation often suggests the acquisition of plasticity and de-differentiation associated with stem cells²⁴ (**Fig. S2A**), together with specific gains in promoter methylation (**Fig. S2B**). To better understand the relationship between genome-wide DNA methylation loss and copy number alterations we examined the correlation between genome-wide DNA methylation frequency and copy number alterations. The Spearman correlation coefficient between methylation frequency and copy number loss was 0.02. The Spearman correlation coefficient between methylation frequency and copy number gain was -0.004 . The lack of correlation between DNA methylation and genomic aberrations in genes for which both methylome and copy number alterations are available in the Discovery cohort can be seen in **Table S5**. The genome-wide DNA methylation and genomic aberrations in tumor samples for which both methylome and copy number alterations are available in the Discovery cohort can be seen in **Figure S3**.

Analysis of Functional Annotation (AFA)²⁵ was then used to integrate the HNSCC methylation, mutation, and expression landscapes and detect alterations in cellular signaling pathways, protein-protein interaction networks, and gene ontology (GO) in HNSCC. AFA revealed that pathways involved in development, differentiation, adhesion, proliferation, and biological/cellular/transcriptional regulation are impacted by concurrent promoter methylation, mutations, and differential gene expression in HNSCC (**Figs. S4 and S5**; **Table S6**). AFA also showed

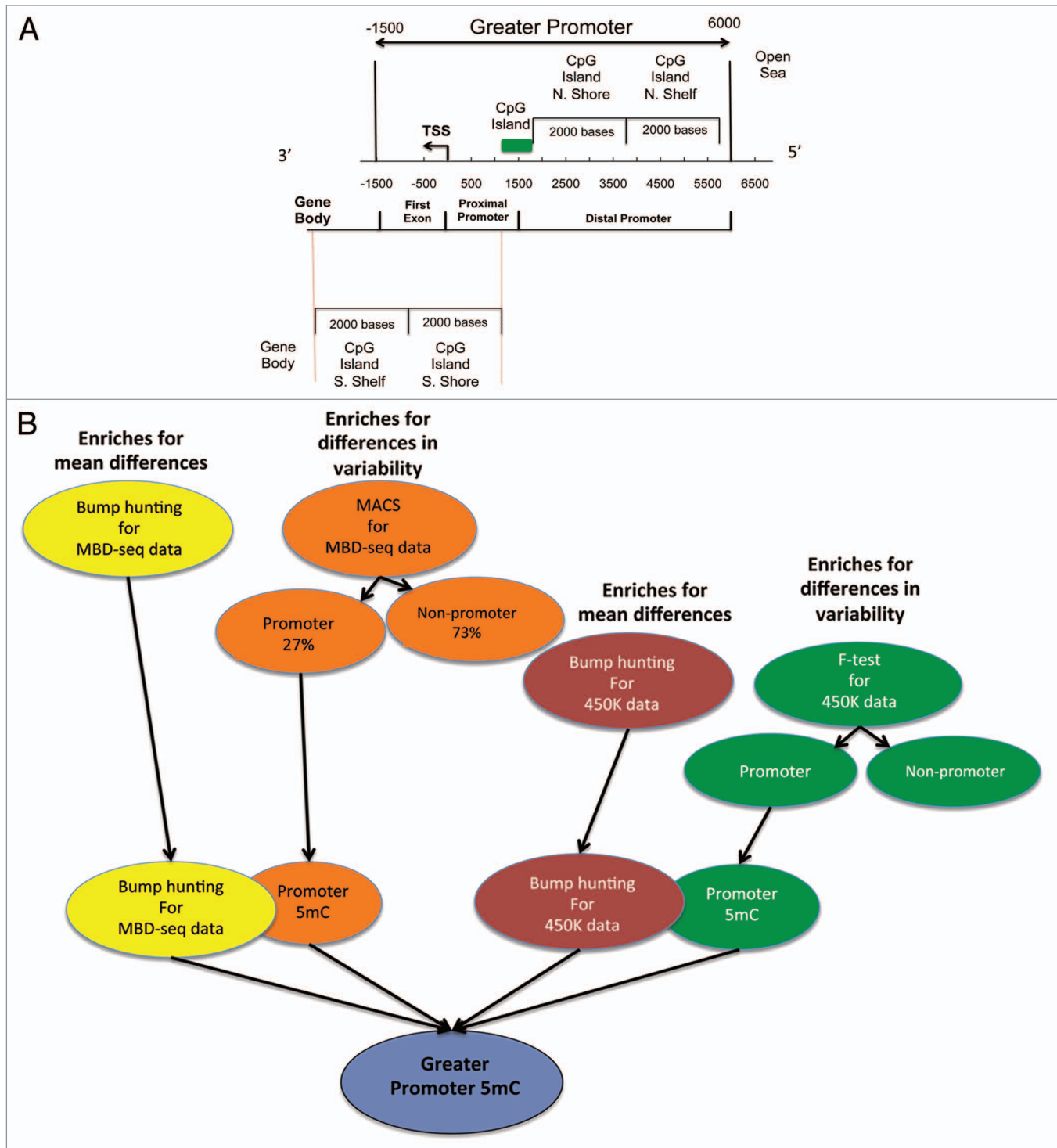


Figure 2. (A) Illustration defining the Greater Promoter region. Using a functional genomic distribution viewpoint we define five CpG genomic locations in relation to their distance to the Transcription Start Site: Proximal promoter, distal promoter, first exon, gene body and intergenic locations. From a CpG content and neighborhood context viewpoint we define four CpG genomic locations in relation to their distance to the nearest CpG Island: CpG Island, CpG Island Shore, CpG Island Shelf, Open Sea and Gene Body. The Greater promoter window is fixed in relation to the TSS. Therefore, the location of CpG Islands will influence the number of significant sequencing reads and 450K probes per gene that are included in our analysis; **(B)** Workflow for identification of differential methylation in the greater promoter of HNSCC, using next-generation MBD-sequencing and 450K methylation platforms. Schematic description of the analytic pipeline developed to unveil the HNSCC methylome. This pipeline enriches for mean genome-wide differences in CpG methylation as also for genome-wide differences in CpG methylation variability at each chromosomal location, for both, methylation sequencing and methylation 450K array data.

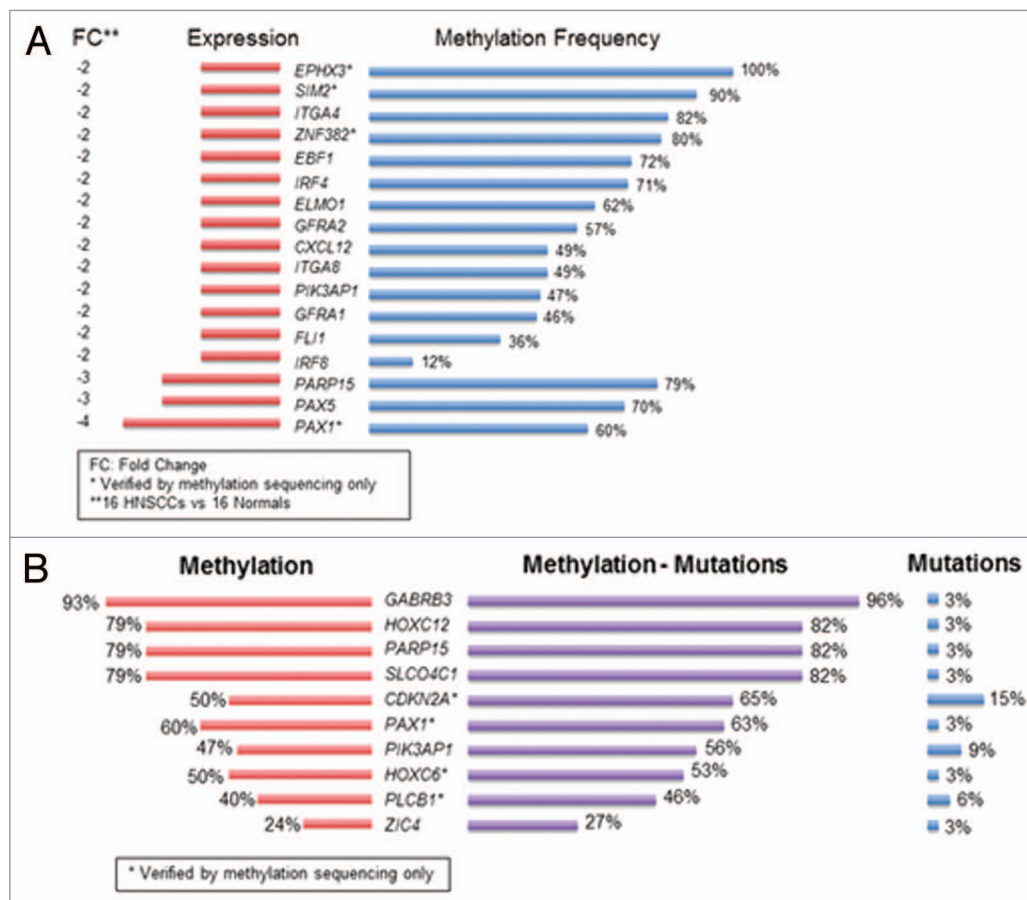


Figure 3. Integrative analysis of co-localized promoter methylation and somatic mutations with concurrent expression changes. (A) Methylated genes with fold change differences in expression greater than 2; (B) Genes with co-localized promoter methylation and somatic mutations in HNSCC. Methylation frequency is represented by the red color. Mutation frequency is represented by the blue color. The purple color represents the combined frequency of methylation and mutation events in the Discovery cohort.

that pathways involved in immune system development, and cell differentiation, proliferation, growth and renewal are impacted by concurrent epigenetic and genetic alterations in HNSCC (Fig. S6).

Methylation-Mutations Interactions with Risk Factors in the Discovery Cohort

When attempting to search for any associations between the mutated and/or methylated genes and HNSCC risk factors (TP53 mutations, HPV status, and smoking history) in the Discovery Cohort, we observed interesting patterns for the *PAX1* and *PAX5* genes. *PAX1* was methylated in all HPV negative tumors whereas zero methylation events of this gene were observed in HPV positive tumors. *PAX1* was also methylated in most patients with a history of tobacco exposure (71%), while only 33% of patients without tobacco exposure history exhibited *PAX1* methylation. Most HPV negative tumors (83%) showed *PAX5* methylation compared with 25% of HPV positive tumors. On the contrary, tumors from patients with a history of tobacco exposure (57%) had similar frequency of *PAX5* methylation to patients with no smoking history (67%). We also observed concurrent genomic and epigenomic associations with viral and tobacco exposures. Patients with *TP53* mutations also had *PAX1*

promoter methylation, history of tobacco exposure, and were HPV negative. Most (83%) of the patients with *TP53* mutations had evidence of *PAX5* methylation (Fig. S7).

Validation of *PAX1*, *ZIC4*, *PLCB1*, and *PAX5* promoter methylation with quantitative methylation specific PCR and TCGA data

We performed qMSP for 3 genes, *PAX1*, *ZIC4*, and *PLCB1*, in 76 tumor samples previously used by Agrawal in a HNSCC deep sequencing study¹ and 19 UPPP samples (Table S7 provides qMSP primers and probes information). All 3 genes appeared on our top ten list of TSG inactivated by both mechanisms and we confirmed a high frequency of tumor specific methylation in the Validation cohort. *PAX1* was methylated in 68% of the cancer cases, *ZIC4* in 80% and *PLCB1* in 52%. We then tested *PAX5* because it is in the *PAX* gene family and appeared on the list of the most frequently inactivated genes by promoter methylation and downregulation of expression. *PAX5* was methylated in 77% of the HNSCC cases in the validation cohort, a number very similar to the 70% methylation frequency identified in the discovery set. We found that *PAX1* ($P < 0.0001$), *ZIC4* ($P < 0.0001$), *PLCB1* ($P < 0.001$), and *PAX5* ($P < 0.0001$) methylation distinguished tumor from UPPP samples (Figs. 4A and B).

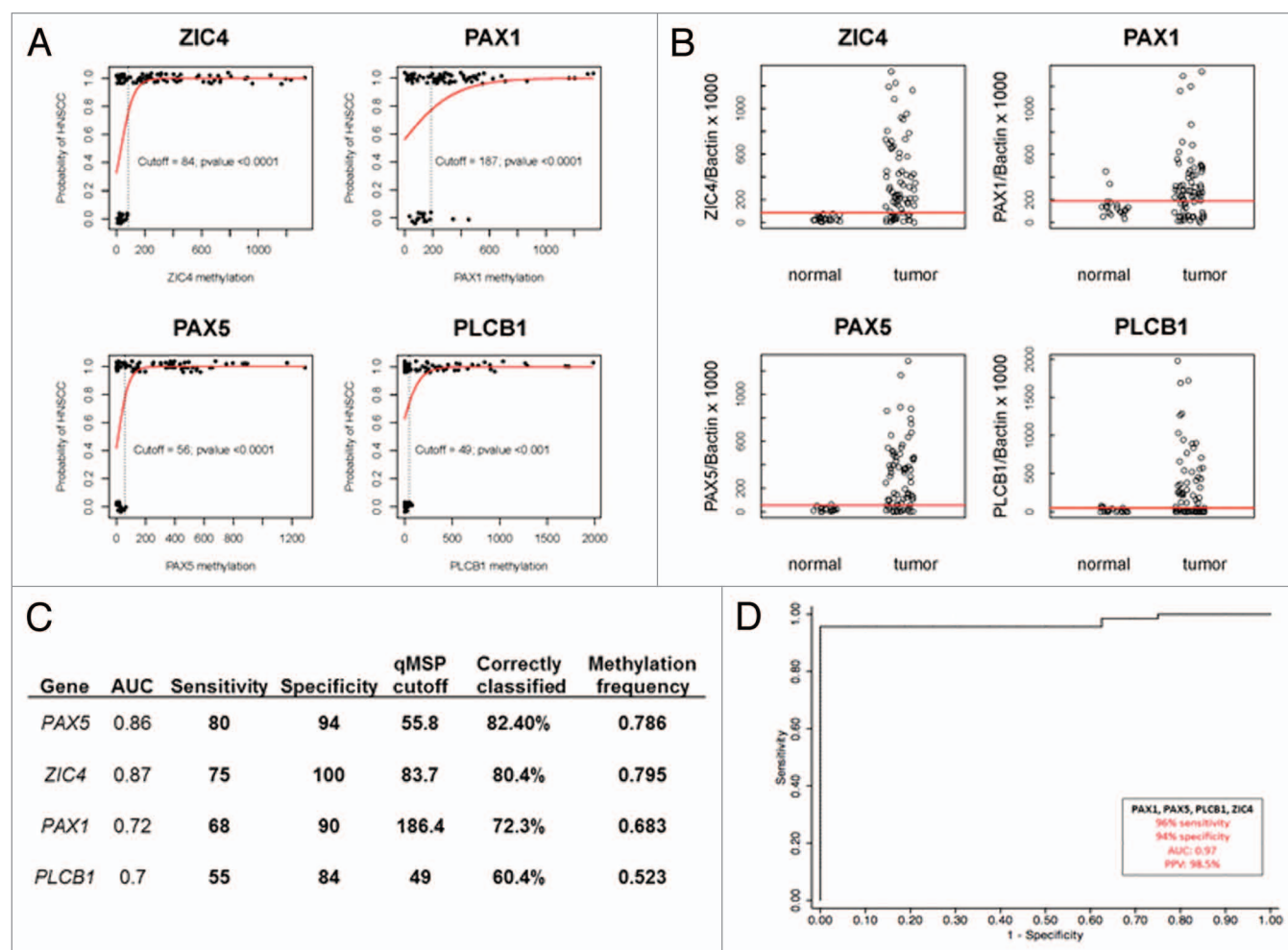


Figure 4. qMSP results for *PAX1*, *PAX5*, *ZIC4* and *PLCB1*. **(A)** Graphical expression of the logistic regression, $\Pr(HNSCC = 1) = \logit^{-1}(b_0 + b_1 \times \text{methylation})$ in tissue from 76 participants with data overlay. The predictor *methylation* is the qMSP value for each case (1) and each control (0). Cutoff methylation values for *PAX1*, *PAX5*, *ZIC4* and *PLCB1* are shown by the vertical dotted line. Probability of HNSCC is shown in red; **(B)** Scatterplots of quantitative MSP analysis of candidate genes promoters in the Validation screen cohort, which consisted of 76 HNSCC tumor tissue samples and 19 normal tissue samples obtained from uvulopharyngopalatoplasty (UPPP) procedures performed in non-cancer patients. The relative level of methylated DNA for each gene in each sample was determined as a ratio of MSP for the amplified gene to ACTB and then multiplied by 1000 [(average value of duplicates of gene of interest / average value of duplicates of ACTB) x 1000] for *PAX1*, *PAX5*, *ZIC4* and *PLCB1*. Red line denotes cutoff value; **(C)** Sensitivity, Specificity and AUC results for qMSP analysis; **(D)** Receiver Operator Characteristics (ROC) curve for promoter methylation of *PAX1*, *PAX5*, *ZIC4* and *PLCB1* genes in the validation cohort. The figure shows that for this four gene panel the qMSP results have 96% sensitivity, 94% specificity, a 0.97 AUC and a Positive Predictive Value of 98.5%.

Receiver Operator Characteristic (ROC) curve analysis revealed that *PAX1* had 68% sensitivity, 90% specificity and a 0.72 AUC; *ZIC4* had 73% sensitivity, 100% specificity, and a 0.87 AUC; *PLCB1* had 55% sensitivity, 84% specificity, and a 0.70 AUC; and *PAX5* had 80% sensitivity, 94% specificity, and a 0.86 AUC (Fig. 4C). A gene panel combining promoter methylation results for these four genes had 96% sensitivity, 94% specificity, a 0.97 AUC, and a Positive Predictive Value of 98.5% (Fig. 4D). A chi-square test of independence revealed an association between methylation in *PLCB1* and tumor site, $P < 0.01$. Tumors of the oral cavity and oropharynx were the most frequently methylated.

All samples harboring *CDKN2A* mutations had *PAX1* methylation ($P < 0.0001$) as did most of *TP53* mutated samples ($P < 0.01$). More than half of *NOTCH1* (61.5%, $P < 0.0001$) mutated

samples also exhibited *PAX1* promoter methylation. All the samples with mutations in *FBXW7* ($P < 0.0001$), and most of the samples with mutations in *TP53* (79%, $P < 0.0001$), and *NOTCH1* (92%, $P < 0.0001$) were methylated in the *PAX5* promoter even after controlling for HPV-status and history of tobacco use (Table S8A).

We further corroborated our qMSP and somatic mutation results by analyzing The Cancer Genome Atlas (TCGA) publicly available data from 279 HNSCC patients (<https://tcga-data.nci.nih.gov>). *PAX5* promoter methylation was associated with *TP53* mutations ($P = 0.02$), while *PAX1* promoter methylation was associated with *NOTCH1* mutations ($P < 0.0001$), even after controlling for HPV-status and tobacco use. This evidence suggests a frequent occurrence of previously unreported interactions

between *PAX1* and *PAX5* promoter methylation and exonic mutations in *NOTCH1* and *TP53* in HNSCC, respectively (Table S8B).

To provide additional evidence of expression downregulation of *PAX1* and *PAX5* in HNSCC we compared mRNA expression in HNSCC and UPPP samples. Differential transcript levels for *PAX1* and *PAX5* were confirmed by quantitative RT-PCR in some of the RNA samples used for microarray analysis and RNA samples from an independent set (Fig. S8A and B). Expression levels were studied in 13 HNSCC and 17 UPPP samples that were readily available. The relative expression levels showed consistency with the results obtained for *PAX1* and *PAX5* in the Discovery cohort with genome-wide methylation and mRNA expression platforms.

We also performed *PAX5* knock-in and knock out studies in *p53* wild type (*p53wt*) and *p53* mutated (*p53mut*) HNSCC cell lines to assess the role of *PAX5* as a tumor suppressor gene connected to the *p53* pathway in HNSCC. We studied the functional consequences of *PAX5* induction in 022(*p53wt*) and 22A(*p53mut*) HNSCC cell lines. Following transfection with (Myc-DDK-tagged)-Human paired box 5, 022 and 22A cells exhibited a dramatic decrease in cell proliferation and *PAX5* expression levels were significantly increased, when measured 48h after transient transfection (Fig. S9).

Conversely, 22B(*p53mut*) HNSCC cells showed modest increase in cell proliferation compared with the control, when *PAX5* was inhibited by siRNA following transfection with *PAX5* siRNAs. Expression levels were significantly decreased, 48h after transient transfection (Fig. S10).

Proposed pathway interplay

After gene set enrichment analysis revealed that concurrent promoter methylation, mutations, and differential gene expression impacted cell differentiation, proliferation, growth and renewal pathways we performed a review study of the most important, potentially impacted cancer pathways in HNSCC. We found evidence of a strong interplay between somatic mutations in *p53* and *NOTCH1* and gene downregulation associated to *PAX1* and *PAX5* promoter methylation in HNSCC.

p53-PAX5

Our literature search revealed that *p53-PAX5* interactions are implicated in apoptotic and/or proliferating signals. *p53* is a downstream target for *PAX* proteins.^{26,27} The human *p53* gene harbors a *PAX* binding site within its un-translated first exon that is conserved throughout evolution, which suggests the importance of this interaction. Frequent promoter methylation of *PAX5* has been reported in ductal carcinoma in situ, invasive breast cancer, and neuroendocrine carcinomas.^{28,29} Furthermore, *PAX5* has been reported to function as a tumor suppressor gene in hepatocellular carcinoma³⁰ and gastric cancer,³¹ directly binding to the *p53* promoter (Fig. S11). *PAX5* also plays an important role in the commitment of lymphoid progenitors to the B lymphocyte lineage.³² The mechanism through which *PAX5* acts in B-cell differentiation is well established.³³⁻³⁵ Recent studies identified some of these interactions also in solid tumors, but little has been shown so far (Fig. 5A).³⁶⁻³⁸ In our Discovery cohort we identified high frequency of *PAX5* promoter methylation in HNSCC,

coinciding with low expression levels, a finding that supports a TSG function.

NOTCH1-PAX1

Agrawal et al. revealed inactivating mutations in *NOTCH1* gene, depicting its importance in HNSCC, proposing also a tumor suppressor function in this particular type of tumors.¹ When *NOTCH1* acts as a TSG it inhibits proliferation and promotes entry to differentiation.³⁹ Targeting the Notch proliferation pathway is really difficult. Preliminary data in cell lines showed that downregulation of *NOTCH1* in *NOTCH1* mutant cell lines leads to a modest effect of cell cycle acceleration and anti-*NOTCH1* agents are only effective in *NOTCH1* wild type cell lines.⁴⁰ In our study, *PAX1* was found to be the most frequently methylated and downregulated gene. In addition to that, several *HOX* family genes, some of which are known to interact with *NOTCH* signaling, were prominent in our list of methylated genes in HNSCC (Fig. S12). Our evidence suggests an interaction between *NOTCH1* and *PAX1* through the *HOX* family of transcription factors^{39,41-43} and also with the Hedgehog pathway through *Hes1*, in which the well-established CCND1 amplification plays an important role.⁴⁴⁻⁴⁸ *PAX1* plays a role in sclerotome differentiation and has been shown to interact with homeobox genes which play a prominent role in normal development and the control of cell proliferation.⁴⁹ Retinoid acid (RA) signaling acts via Hox gene pathways,^{50,51} some of which are able to regulate *PAX1* through canonical *NOTCH1* expression. These interactions are described in Figure 5B.

Discussion

We have conducted the first comprehensive integrated genomic and epigenomic analysis in HNSCC, focusing on identifying TSG genes that demonstrate concurrent promoter methylation with downregulation of expression and somatic mutations. Recent studies in HNSCC, published as we were finalizing this manuscript, focused on therapeutic pathways affected by somatic mutations and copy number alterations,⁵²⁻⁵⁴ but only described the clustering effects of DNA methylation at a global genome wide level.⁵² We performed the first detailed genome wide analysis of the HNSCC methylome that studied expression alterations associated with differential methylation patterns in the greater promoter region, focusing on key TSGs that are also inactivated by somatic mutations. The main focus of this paper was to identify the number of tumor suppressor genes differentially methylated in the greater promoter region and mutated in HNSCC, and examine their combined impact in expression downregulation. We controlled for chromosomal deletions using CNV data previously generated in the Discovery cohort, to distinguish if expression alterations were related to methylation events or to deletions in specific regions of the chromosome.

Many genes with dense promoter methylation and downregulation of expression are candidate TSGs. However, genes inactivated by both somatic mutations and promoter methylation are likely to be key drivers of oncogenesis. Our analysis identified 10 downregulated genes inactivated by somatic mutation and

promoter methylation. We selected *PAX1*, *ZIC4*, *PLCB1*, and *PAX5* for further validation and we were indeed able to detect stark differences in DNA methylation levels between cancer and UPPP samples. *ZIC4* and *PLCB1* were the genes with the lowest methylation frequency in HNSCC, exhibiting also low mutation frequency, and yet, we confirmed *ZIC4* and *PLCB1* promoter methylation differences between normal and cancer samples in the Validation cohort.

PAX1 and *PAX5* were highlighted as key genes in this study as they belong to the same family of transcription factors and were both found to be methylated and downregulated in HNSCC. *PAX1* and *PAX5* are genes involved in differentiation/proliferation signals and the gene set enrichment analysis performed clearly depicted these signaling pathways as deregulated in HNSCC in our discovery set of tumors.

We observed interesting relationships among the most commonly mutated and methylated genes in HNSCC: 61.5% of the *NOTCH1* mutated samples also exhibited *PAX1* methylation and 79% of the samples carrying *TP53* mutations were also methylated in the *PAX5* gene promoter. External validation in 279 primary HNSCC samples from the TCGA project verified our initial findings. Combined, this evidence suggests the frequent occurrence of previously unreported interactions between *PAX1* and *PAX5* promoter methylation and exonic mutations in *NOTCH1* and *TP53* in HNSCC.

The greater promoter *PAX1* and *PAX5* methylation levels reported in this manuscript were obtained with three different platforms that use diverse technology, chemistry, sample preparation and data analysis pipelines to obtain methylation values. We have combined these three platforms into a very robust methylation detection pipeline.

PAX1 and *PAX5* methylation levels listed in the Discovery cohort are both, from MBD-seq and the 450K BeadChip assay. The 450K results were used to confirm the MBD-seq results in the subset of ten Discovery samples that were sequenced and, by proxy, validate the methylation results of the other 22 Discovery cohort samples that were only queried with the 450K arrays. The levels of *PAX1* and *PAX5* methylation levels listed for the Prevalence cohort were obtained with quantitative methylation specific PCR (qMSP). The levels of *PAX1* and *PAX5* listed for TCGA were obtained with the 450K BeadChip assay. The difference in *PAX1* methylation between HPV positive and HPV negative tumors was significant in the Discovery cohort. The difference between the two in the Prevalence cohort was not significant ($P = 0.225$), perhaps due to the small sample size of HPV positive patients.

PAX genes, a family of nine transcription factors which act as cell lineage specific regulators of the tissues where they are normally expressed, are now also recognized as important factors in cancer progression. *PAX* genes, similarly to the *NOTCH* gene family, may play previously unrecognized fundamental roles in balancing proliferation and differentiation signals, two conceptually opposite cellular processes in canonical cancer research. The *PAX* family of genes may ultimately, following Waddington's epigenetic landscape metaphor, be part of an epigenomic mediated switch between cancer initiation and cancer maintenance

pathways, which stochastically drive cancer progression, immune system avoidance, acquisition of tumor resistance, and establishment of metastatic disease. Loss of *NOTCH1* function due to mutation, or mutation/methylation-dependent silencing of downstream genes, such as *PAX1* or the *HOX* family genes, is likely to abrogate normal cell differentiation.

Clinically, *TP53* mutation has been shown time and again to be among the worst molecular alterations in patients with HNSCC.^{36,55} Patients that harbor p53 mutant tumors are more likely to relapse after complete resection and radiation therapy.⁵ We confirmed a high frequency of *PAX* methylated tumors in 279 HNSCC tumor samples from the TCGA cohort and found that tumors which already harbor a p53 mutation, also harbor *PAX5* promoter methylation.

Together, our results support the notion that differential promoter methylation and somatic mutations are the main cause of gene inactivation and pathway deregulation in HNSCC. We have unveiled hitherto unknown interactions between mutated and methylated genes that are associated with gene expression alterations in HNSCC. Characterization of the complete HNSCC methylome has contributed insights into the clustering of specific genetic and epigenetic events in the greater promoter region and highlights the importance of understanding the relative contribution of each to the overall frequency of TSG inactivation. We plan future in vitro studies focused on the functional consequences of the inactivation of these high frequency genes and will further explore the above proposed gene interactions. Understanding the complete contribution of genomic and epigenomic alterations to specific genes and pathways in cancer will reveal novel high frequency specific markers for better risk assessment (as we observed in the TCGA cohort above) and will highlight the true frequency of therapeutic pathways to better target the disease at the molecular level.

Materials and Methods

Participants

Patient selection

Head and Neck Squamous Cell Carcinoma ($n = 91$) and uvulopalatopharyngealplasty (UPPP) patients ($n = 35$) were consented for this study at the Johns Hopkins Medical Institutions hospitals and MD Anderson Cancer Center. The study was approved by the Ethics Committee of each participating hospital, as well as by the Johns Hopkins Institutional Review Board.

Johns Hopkins component

Fresh-frozen surgically resected tissue and matched blood were obtained from patients at Johns Hopkins Medical Institutions in Baltimore. Tissue was analyzed by frozen section histology to estimate neoplastic cellularity. In order to enrich the samples for neoplastic cells, normal tissue was removed from the samples using macro-dissection based on the frozen section histology. HPV tumor status was determined for oropharyngeal tumors per standard clinical care using in situ hybridization. Hybridization was performed using the HPV III Family16 probe set that captures HPV genotypes 16, 18, 33, 35, 45, 51, 52, 56, and 66.

HPV16-positive controls included an HPV16-positive oropharyngeal cancer and the SiHa and CaSki cell lines. HPV tumor status was also determined by E6/E7 PCR primer amplification.

MD Anderson Component

Fresh-frozen surgically resected tumor and matched non-malignant adjacent tissue were obtained from consented patients treated for HNSCC at the University of Texas MD Anderson Cancer Center, under an Institutional Review Board approved protocol. Frozen tissue was embedded in optimal cutting temperature compound and cryosections from the top and middle of specimens were stained with hematoxylin and eosin prior to being evaluated by a pathologist for the presence of >60% tumor nuclei content or absence of tumor (i.e., normal). Samples that passed this criterion were sectioned all the way through and washed once in PBS prior to isolating genomic DNA using an ArchivePure DNA purification kit.

Methylated binding domain sequencing (MBD-seq)

Preparation of libraries

Tissue samples were digested with 1% SDS and 50 µg/mL proteinase K (Boehringer Mannheim) at 48 °C overnight, followed by phenol/chloroform extraction and ethanol precipitation of DNA. Two micrograms of DNA were sonicated to a modal size of ~150–250 bp, and end-repaired using the NEBNext SOLiD DNA library preparation kit end-repair module following the manufacturer's protocol (New England Biolabs). After column-purification (using the Qiagen PCR purification kit), SOLiD P1 and P2 adapters lacking 5' phosphate groups (Life Technologies) were ligated using the NEBNext adaptor ligation module and column-purified, and subjected to nick-translation by treating with Platinum Taq polymerase to remove the nick.

Affinity enrichment and capture of methylated DNA fragments

The resulting library was divided into two fractions, a total input fraction, and an enriched methylated fraction. The enriched methylated fraction was then subjected to affinity enrichment of methylated DNA fragments by using 6xHis-MBD2-MBD polypeptides immobilized on magnetic beads as described previously.^{56,57} The resulting enriched methylated fraction and the total input fraction were then subjected to library amplification using the NEBNext amplification module according to the manufacturer's protocols, using 4–6 cycles for the total input, and 10–12 cycles for the enriched methylated fractions. Library fragments that were between 200–300 bp were size selected after agarose gel electrophoresis.

Massively parallel sequencing of MBD-seq libraries

The libraries were then subjected to emulsion PCR and bead enrichment following the SOLiD emulsion PCR protocol (Life Technologies). The resulting beads were then deposited on the SOLiD flow cell and subjected to massively parallel 50 bp single-read sequencing on a SOLiD v4.0 sequencer octet segment, with one octet segment for the total input and another one for the enriched methylated fraction. The details of the sequencing output for each sample from the sequencing run (number of tags, coverage, etc.) are provided in Table S9. Reads were aligned to hg19 using default settings in bioscope v1.3, with the exception of the bam output method, which was changed to alignment score.

Bioinformatics analysis of MBD-seq data

MACS analysis and identification of differential methylation

For the purposes of the analysis we divided the genome into two broad regions: the greater promoter, was defined as the region encompassing 6000 bases upstream and 1500 bases downstream from the transcription start site (TSS). From the functional genome distribution standpoint the greater promoter region, includes CpG sites in the proximal promoters, 1500 bases upstream from the described TSS, and 1500 bases downstream from the TSS, in the 5' untranslated region and exon1. From the CpG content and neighborhood context the differentially methylated CpGs in HNSCC were located in CpG islands, CpG shores (regions 2000 bp upstream and downstream of but not inside CpG islands), CpG shelves (regions 2000 bp upstream and downstream of but not inside the shores), or as isolated CpGs in the area of the genome now defined as Open Sea. Methylated regions were identified as peaks of aligned sequencing tags in the enriched compared with total input fraction using MACS v1.4 software,^{22,58,59} which allows identification of peaks after accounting for both global and local biases using the total input fraction. Peaks of methylation were identified for each sample separately.

To identify differentially methylated regions we first used stringent parameters to define presence and absence of methylation in EACH sample as follows: we used a low MACS *P* value cut-off ($P < 10^{-6}$) to identify regions that are methylated, and another cut-off (MACS $P > 10^{-2}$) to identify those regions that have very little evidence for methylation.

Next, for any given comparison of group A vs. group B (e.g., Group A = all tumors; Group B = all normals), we identified all regions that showed absence of methylation in all samples of Group A and presence of methylation in at least one sample from group B. All such overlapping regions across samples with peak calls in Group B were then merged, and the number of such samples and the lowest p-value of the peaks for these samples were recorded as the aggregate differentially methylated region. This analysis therefore yields regions in which all samples in Group A have absence of methylation peaks, and at least one sample in Group B has a methylation peak. The converse comparisons (i.e., absence of methylation in Group B, with at least one sample having a methylation peak in Group A) were also performed to obtain regions showing both gain of- and loss of-methylation.

Methylation bump hunting for identification of differentially methylated regions

Differential methylation was also identified using an independent approach called bump hunting that has been previously used to identify differential peaks in methylation data. Methylation bump hunting is a data analysis pipeline that effectively models measurement error, removes batch effects, detects regions of interest and attaches statistical uncertainty to regions identified as differentially methylated.²³ These methods are implemented in the *bumphunter* Bioconductor package and described in more detail in the section "Bioinformatics for 450K data." Reported functionally relevant findings have been generally associated with genomic regions rather than single CpGs, either CpG islands,⁶⁰ CpG island shores,²⁰ genomic blocks,⁶¹ or generic 2-kb regions.⁶²

Epigenomic bumps may have greater variability in size and shape than MBD-seq peaks.

Integration of MACS and bumphunting results

To identify the promoter regions that are differentially methylated in HNSCC when compared with normal oral mucosa, we intersected the list of methylated probes that discriminated between tumor and normal tissue identified with MACS with the list of methylated regions that discriminated between tumor and normal tissue identified with bumphunting. We used R (v3.00) to analyze the correlation of methylated promoter regions and HNSCC etiological factors.

Verification of MBD-seq results with HumanMethylation450K DNA BeadChip assay

450K array description and sample preparation

The 450K is a two-color array that detects cytosine methylation at 485,512 methylation loci, mostly at CpGs, but also at a small number of cytosine residues outside of the CpG context, using bisulfite converted DNA. For each individual CpG two different signals are detected. One signal measures the amount of methylated DNA (Meth) and the other one measures the amount of unmethylated DNA (Unmeth). The Meth and Unmeth signals are measured with two different assays called “Type I” design or a “Type II” design. A β (β) value is generated by both Type I and Type II design probes to denote the methylation level of the CpG loci using the ratio of intensities between Meth and Unmeth (β value = methylation intensity / methylation + unmethylated intensity of the given CpG locus). Each methylation locus is interrogated by one of these designs. For a type I locus the Meth and Unmeth signals are measured by two paired probes, with a given locus using either the red or green signal from these probes. Type II loci are assayed using a single probe, with Meth and Unmeth signals derived from the green and red channels respectively. In addition to the methylation loci, the array contains a small number of control probes and 65 probes measuring common SNPs, intended for sample tracking. Type II probes use only one probe per methylation locus and hence allows more loci on the array, at a fixed array size. However, due to the chemistry used by the type II probe design, type II probes can only tolerate up to three CpGs within the 50bp probe. The type I design tolerates more CpGs within the 50bp probe, but assumes that all methylation loci in the probed sequence are in the same state.⁶³

Bisulfite modification of genomic DNA (2 μ g) was performed with EpiTect Bisulfite Kit (QIAGEN) according to the manufacturer’s protocol. We hybridized bisulfite converted DNA from normal (UPPP) tissue ($n = 16$) and Head and Neck Squamous Cell Carcinoma (HNSCC) tissue ($n = 31$) samples to the 450K array.

Bioinformatics for 450K data

Bioinformatics strategies were used for background correction, normalization, and data analysis of differentially methylated genomic regions between tumor, and normal tissue. As with the analysis of methylation sequencing data we used two different analytic pipelines: a pipeline designed to capture the variability in methylated signals across the arrays using an F-test¹¹ and the bumphunting pipeline.²³ We used the *minfi* and *bumphunter* packages found in Bioconductor to perform background

correction, normalization, and data analysis of differentially methylated genomic regions between tumor and normal tissue. The *minfi* package provides tools for analyzing Illumina’s Methylation arrays, with a special focus on the new 450k array for humans, and includes methods for preprocessing, quality assessment, and detection of differentially methylated regions from the kilobase to the megabase scale.⁶⁴ The *bumphunter* package is meant to work on data with several biological replicates, similar to the *lmFit* function in *limma*. While *bumphunter* is written using genomic data as an illustrative example, most of it is generalizable to other data types (with some one-dimensional location information).

F-test analytic pipeline

The selection of significantly methylated CpGs in the Illumina 450K Infinium assay data was performed in a stepwise manner. The 450.idat files were preprocessed and background corrected with the *preprocessIllumina* function in *minfi* to obtain β values. An F-test was performed across all 47 samples to identify CpGs with a significant difference in β values between normal and malignant tissue. Since the empirical P values were calculated genome-wide, adjustment for multiple testing was performed. Q-values were computed from the empirical P values using the Benjamin and Hochberg correction. Probes with q-values less than 0.05 were deemed statistically significant and were included in the CpG list. We then selected only those CpGs that showed a methylation difference of at least 0.25 between cancer and normal tissues and a β value of at least 0.3 in cancer, as previously described.¹¹ All bioinformatics analyses were performed using R version 3.0.

450K-bumphunting analytic pipeline

We performed pre-processing with the *minfi* package, which applies a version of subset quantile normalization to the Meth and Unmeth intensities separately. The distribution of type I and type II probes is forced to be the same by first quantile normalizing the type II probes across samples and then interpolating a reference distribution to which the type I probes are normalized. For the probes on the X and Y chromosomes the males and females are normalized separately. Sex is determined by the *getSex* function using copy number information. The stratified quantile normalization method is implemented by the *preprocessQuantile* function (the function does no background correction and removes zeros using the *fix2MethOutlier* function). This algorithm relies on the assumptions necessary for quantile normalization and involves both within- and between- sample normalization.⁶⁴

Integration of F-test and 450K-bumphunting results

To identify the promoter regions that are differentially methylated in HNSCC when compared with normal oral mucosa, we intersected the list of methylated probes that discriminated between tumor and normal tissue identified with the F-test with the list of methylated regions that discriminated between tumor and normal tissue identified with 450K-bumphunting.

E. mRNA expression arrays

Total RNA was isolated from normal (UPPP) tissue ($n = 16$) and Head and Neck Squamous Cell Carcinoma (HNSCC) tissue ($n = 16$) samples by using Tri-reagent. cDNA was made, and

hybridized to Affymetrix GeneST1.0 Arrays (Affymetrix) according to manufacturer's instructions. Six of these samples were also sequenced with MDB-seq. The data obtained from CEL files was background corrected with RMA, quantile normalized before an ANOVA was used to determine the Fold Change difference in log-transformed intensities between Tumor and Normal samples.

Analysis of functional annotation

Enrichment analysis of functional themes (Analysis of Functional Annotation, AFA) was performed to capture biological processes over-represented in the various conditions under investigation. This unbiased computational approach, conceptually similar to Gene Set Enrichment Analysis (GSEA),⁶⁵ enables the interpretation of genome-wide data through the identification and visualization of information encompassing distinct biological concepts, and was previously used successfully to integrate and interpret both differential gene expression and methylation data.^{25,66} A chi-square test of independence was applied to test whether each Functional Gene Set (FGS) was over-represented in any of the gene list associated with any of the investigated contrasts/conditions (e.g., gene associated with methylated promoters in HNSCC). In the present study, individual, non-redundant genes, as annotated in the NCBI Entrez gene database (R/Bioconductor package org.Hs.eg.db version 2.4.6) were used as the total gene space, and contingency tables were used to identify gene sets over-represented in the investigated conditions. Correction for multiple hypothesis testing was obtained separately for each FGS collection, by applying the Benjamini and Hochberg method⁶⁷ as implemented in the multi-test R/Bioconductor package. Overall, this approach is analogous to Gene Set Enrichment Analysis (GSEA),^{65,68,69} and has already been successfully applied in other studies.^{66,70} The heat-maps' color bar represents the negative log₁₀ False Discovery Rate (FDR). For each gene set collection the sets for which at least one condition showed FDR < 0.01 were reported. The top 150 conditions were reported when too many gene sets were retrieved.

AFA was used to compare biological themes enriched in the following gene lists: (1) mutated genes; (2) genes with hyper-methylated promoters; (3) genes showing hyper-methylation outside the greater promoter region; (4) genes upregulated in cancer when compared with normal; and (5) genes downregulated in cancer when compared with normal. Gene set enrichment was assessed by testing for gene set over-representation with a chi-square test, because each gene list was obtained from diverse analyses using different methods. In the AFA analysis each gene was counted only once, while using the totality of the genes annotated to the NCBI Entrez gene database as the background gene space. For this reason there was no confounding with the number of probes or the length of the genes (i.e., each gene was counted only once irrespective to the number of probes or its length). The GO categories reported in Table S6 encompass different biological processes and contain a fairly large number of distinct genes. Such genes show variable length, from short to long transcripts, and all possible and disparate genomic arrangements, with genes located in both "gene-rich" and "gene-poor" genomic regions.

Validation of genomic and epigenomic alterations

Quantitative methylation specific PCR (qMSP) in the prevalence cohort

qMSP was used for validation in a Prevalence cohort of 76 tumors in which we had previously identified somatic mutations in *TP53*, *NOTCH1*, *CDKN2A*, *PIK3CA*, *FBXW7*, and *HRAS* and in 19 UPPP normal control tissue samples. We examined promoter methylation in three of the genes that were included in our final list of mutated and methylated genes and that were methylated in at least 40% of the Discovery Cohort samples: *PAX1*, *PAX5*, *PLCB1*, and *ZIC4*. Bisulfite-modified DNA was used as a template for fluorescence-based real-time PCR, as previously described.⁷¹

Contingency tables of mutational and methylation events in TCGA data set

Publicly available HNSCC Illumina 450K methylation and exome sequencing data was downloaded from the TCGA website (<http://cancergenome.nih.gov>) and the cBioPortal for Cancer Genomics (www.cbioportal.org/public-portal/) using R (v3.0.0). Publicly available exome mutation data for *TP53* and *NOTCH1* and β values for all *PAX1* and *PAX5* 450K array probes were extracted for all HNSCC samples analyzed by the TCGA project that had paired methylation and mutation data for the genes of interest ($n = 279$). Only the 450K probes located in TSS1500, TSS200, and 1st exon, as per the manufacturer's annotation, were used to create the contingency tables for methylation and mutation analyses. Contingency tables were used to examine the association between exonic mutations of *TP53*, *CDKN2A*, *HRAS*, *FBXW7*, and *NOTCH1* and promoter methylation of *ZIC4*, *PAX1*, *PAX5*, and *PLCB1*. The MacNemar test for paired data was implemented in R (v3.0.0) to evaluate the association between mutations and promoter methylation.

Quantitative real-time reverse transcription PCR

HNSCC RNA samples from the Discovery cohort and from an independent cohort were assessed for *PAX5*, *PAX1*, and *GAPDH* expression levels using quantitative real-time reverse transcription (RT)-PCR (TaqMan). Reverse transcription was performed with random hexamer primers and Superscript II Reverse Transcriptase (Invitrogen Corp.) according to manufacturer's instructions. Quantitative RT-PCR was then performed on the Applied Biosystems 7900 Sequence Detection Instrument (Applied Biosystems) using TaqMan expression assays (Life Technologies).

Functional studies in cell lines

Human HNSCC cell lines with known *p53* status were cultured to determine *PAX5* methylation and expression levels. Human HNSCC cell lines 022 (*p53* wt), 22A (*p53* mut), and 22B (*p53* mut) were selected for functional studies based on their expression and methylation results (Fig. S13). Cell growth conditions were maintained at 37 °C in an atmosphere of 5% CO₂.

Transient transfection, PAX5 inhibition or overexpression, and cell proliferation assay

We knocked down *PAX5* in the 22B cell line using the ON-TARGETplus Pool of siRNAs against *PAX5* and a non-targeting Pool of siRNAs (Thermo Scientific) as control. Cells were seeded in 96-well plates and allowed to grow until approximately

70% confluence. Cells were transfected with siRNA using Lipofectamine RNAiMAX Reagent (Invitrogen) and cell metabolic activity was determined every 24 h using the CCK-8 colorimetric assay (Dojindo). Values are mean \pm SEM for pentaplicates of cultured cells. The transfection efficiency was confirmed by quantitative real-time RT-PCR as described above and normalized to *GAPDH* at 48-h time point. Pax5 knockdown in 22B cells showed a modest increase in cell proliferation compared with the control.

Forced expression of *PAX5* in 022 and 22A cell lines was performed. The pCMV-Entry-EV (control) and the (Myc-DDK-tagged)-Human paired box 5 (*PAX5*) from ORIGENE were transfected into cells (FUGENE HD Promega) which were then seeded in 96-well plates and allowed to grow until approximately 70% confluence. Cell metabolic activity was determined every 24 h using the CCK-8 colorimetric assay (Dojindo). Values are mean \pm SEM for pentaplicates of cultured cells. The transfection efficiency was confirmed by quantitative real-time RT-PCR as described above and normalized to *GAPDH* at the 48-h time point. 022 and 22A cells exhibited a dramatic decrease in cell proliferation after forced expression of *PAX5* 48h after transfection compared to controls.

Disclosure of Potential Conflicts of Interest

J.A.C. is the Director of Research of the Milton J. Dance Head and Neck Endowment. V.E.V. is a co-founder of Inostics

and Personal Genome Diagnostics and is a member of their Scientific Advisory Boards. V.E.V. owns Inostics and Personal Genome Diagnostics stock, which is subject to certain restrictions under University policy. The Johns Hopkins University in accordance with its conflict of interest policies is managing the terms of this arrangement.

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Supplemental Materials

Supplemental materials may be found here: www.landesbioscience.com/journals/epigenetics/article/29025

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